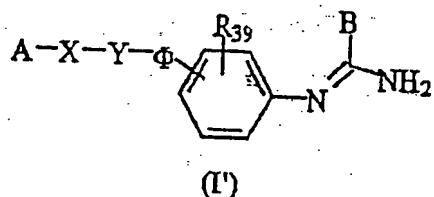


In the Claims:

Claims 1 to 13 (cancelled).

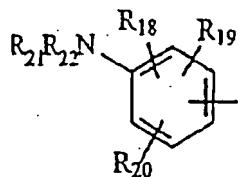
Claim 14 (currently amended)

A compound of general the formula (I')



wherein

A is



R_{18} , R_{19} and R_{20} are independently selected from the group consisting of hydrogen, halogen, -OH, SR_{23} , alkyl or alkoxy of 1 to 6 carbon atoms, alkenyl of up to 6 carbon atoms and $\text{NR}_{24}\text{R}_{25}$; R_{21} and R_{22} are independently selected from the group consisting of hydrogen and alkyl of 1 to 6 carbon atoms, or R_{21} and R_{22} form together with the nitrogen atom an optionally substituted heterocycle having 4 to 7 members and 1 to 3 heteroatoms including the already present nitrogen atom, the additional heteroatoms being independently selected from the group consisting of O, N or furthermore R_{21} is selected

from the group consisting of alkylsulfonyl, alkylsulfoxide and alkylcarbonyl and then R₂₂ is hydrogen, R₂₃ is hydrogen or alkyl of 1 to 6 carbon atoms, R₂₄ and R₂₅ are independently selected from the group consisting of hydrogen, OH, alkyl of 1 to 6 carbon atoms and CO, R₂₆, R₂₆ is alkyl of 1 to 6 carbon atoms,

B is selected from the group consisting of alkyl of 1 to 6 carbon atoms, NR₃₄R₃₅, carboyclic or heterocyclic aryl with 5 or 6 members containing from 1 to 4 heteroatoms selected from the group consisting of O, S and N, the aryl radical being optionally substituted by at least one member selected from the group consisting of alkyl or alkoxy of 1 to 6 carbon atoms and alkenyl of up to 6 carbon atoms thiophenyl,

R₃₄ and R₃₅ are independently selected from the group consisting of hydrogen and alkyl of 1 to 6 carbon atoms, or R₃₄ and R₃₅ form together with the nitrogen atom a non-aromatic heterocycle with five to six members, each of the elements of the chain being selected from the group consisting of CH₂, NH, O and S,

X is selected from the group consisting of a bond, -(CH₂)_m, -(CH₂)_mCO, O(CH₂)_m, -S(CH₂)_m, NR₃₆(CH₂)_m, or -CO-NR₃₆-, -O(CH₂)_mCO, S(CH₂)_mCO, NR₃₆-(CH₂)_mCO, (CH₂)_mC(OH)(CH₃)CO, CH=CH and CH=N,

Y is selected from the group consisting of a bond, -(CH₂)_n, and -(CH₂)_rQ-(CH₂)_s-, and thiazolidine,

~~Q is selected from the group consisting of piperazine, homopiperazine, 2-methylpiperazine, 2,5-dimethylpiperazine, piperidine, 1,2,3,6-tetrahydropyridine, pyrrolidine, azetidine, thiazolidine and a saturated carbon ring having 3 to 7 members,~~

~~Φ is selected from the group consisting of a bond, $(CH_2)_p-O-(CH_2)_q$, $(CH_2)_p-S-(CH_2)_q$, $-(CH_2)_p-NR_{37}-(CH_2)_q$, $-(CH_2)_p-CO-NR_{37}-(CH_2)_q$, and $CO(CH_2)_p-NR_{37}-(CH_2)_q$,~~

~~R₃₆ and R₃₇ are independently selected from the group consisting of hydrogen, alkyl of 1 to 6 carbon atoms and -CO-R₃₈, R₃₈ is alkyl or alkoxy of 1 to 6 carbon atoms,~~

~~R₃₉ is selected from the group consisting of hydrogen and alkyl or alkoxy of 1 to 6 carbon atoms,~~

~~m, n, p, q, r and s are independently integers from 0 to 6,~~

~~and its pharmaceutically acceptable salts.~~

Cancel Claims 15 to 19.

Claim 20 (previously presented) A compound of claim 14 selected from the group consisting of

~~- 2-amino-N-(4-([amino(2-thienyl)methylidene]amino)phenethyl)-5-methoxybenzamide;~~

- 5-amino-N-(4-{{[amino(2-thienyl)methylidene]amino}phenethyl}-2-hydroxybenzamide;
- 4-(4-{{[amino(2-thienyl)methylidene]amino}phenyl}-N-{{[methylsulphonyl]amino}phenyl}butanamide;
- 4-(4-{{[amino(2-thienyl)methylidene]amino}phenyl}-N-[4-(dimethylamino)phenyl]butanamide;
- 5-(4-{{[amino(2-thienyl)methylidene]amino}phenyl}-N-[4-(dimethylamino)phenyl]pentanamide;
- (4*R*)-2-(3-{{[amino(2-thienyl)methylidene]amino}-phenyl}-N-[4-(dimethylamino)phenyl]-1,3-thiazolidine-4-carboxamide;
- *tert*-butyl 3-{{[amino(2-thienyl)methylidene]amino}benzyl}{3-[4-(dimethylamino)anilino]-3-oxopropyl}carbamate;
- 3-{{(3-{{[amino(2-thienyl)methylidene]amino}-benzyl)amino}-N-[4-(4-methyl-1-piperazinyl)phenyl]propanamide;
- 3-{{(3-{{[amino(2-thienyl)methylidene]amino}-benzyl)amino}-N-[4-(4-morpholinyl)phenyl]propanamide;
- N'-[4-(2-{{[5-(dimethylamino)-2-hydroxybenzyl]amino}ethyl)phenyl]-2-thiophenecarboximidamide;
- N-{{(4-{{(4-{{[amino(2-thienyl)methylidene]amino}phenethyl)-amino}methyl}phenyl)acetamide;
- N'-[4-(2-{{[5-(dimethylamino)-2-hydroxy-3-methoxybenzyl]amino}ethyl)phenyl]-2-thiophenecarboximidamide;
- N'-{4-[2-({[4-(dimethylamino)anilino]carbonyl}amino)-ethyl]phenyl}-2-thiophenecarboximidamide;
- N'-{4-[2-{{[5-(dimethylamino)-2-hydroxy-3-methoxybenzyl]-(methyl)amino}ethyl}phenyl]-2-thiophenecarboximidamide;

and the pharmaceutically acceptable salts of the latter.

Claim 21 (withdrawn) A method of inhibiting NO synthase in a patient in need thereof comprising administering to said patient a therapeutically effective amount of a compound of claim 14.

Claim 22 (withdrawn) A method of inhibiting lipidic peroxidation in a patient in need thereof comprising administrating to said patient a therapeutically effective amount of a compound of claim 14.

Cancel Claim 23.

Claim 24 (withdrawn) A method of treating a neurodegenerative disease in a patient in need thereof comprising administrating to said patient a therapeutically effective amount of a compound of claim 14.

Claim 25 (withdrawn) The method of claim 24 wherein the neuorodegenerative disease is selected from the group consisting of Alzheimer's disease, Huntington's chorea, Parkinson's disease, Creutzfeld Jacob disease and amyotrophic lateral sclerosis.